This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A compound of Compounds of the formula I

in which

R denotes Hal, -C≡C-H, -C≡C-A or OA,

R¹ denotes H, =O, Hal, A, OH, OA, A-COO-, Ph-(CH₂)_n-COO-, cycloalkyl-(CH₂)_n-COO-, A-CONH-, A-CONA-, Ph-CONA-, N₃, NH₂, NO₂, CN, COOH, COOA, CONH₂, CONHA, CON(A)₂, O-allyl, O-propargyl, O-benzyl, =N-OH, =N-OA or =CF₂,

X, X' each, independently of one another, denote CH, CHal or N,

Y denotes R⁴ or Hal,

Ph denotes phenyl which is unsubstituted or mono-, di- or trisubstituted by A, OA, OH or Hal,

R² denotes H, Hal or A,

R³ denotes H or A.

R⁴ denotes OH, OA, A-COO-, NHA, NHAr, NAA', Het or -NH-CHR⁵-COOR³,

R⁵ denotes H, A, -CHR³-OH, $(CH_2)_n$ -Ph, $(CH_2)_n$ -COOH, $(CH_2)_n$ -CONH₂, $(CH_2)_n$ -NH₂, $(CH_2)_n$ -NH(=NH)NH₂, $(CH_2)_n$ -Het¹ or $(CH_2)_n$ -SR³,

Het denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be unsubstituted or mono-, di- or trisubstituted by A, OH, OA, CN, COOH, COOA and/or carbonyl oxygen (=O),

Het¹ denotes a mono- or bicyclic aromatic heterocycle having 1 to 4

N, O and/or S atoms, which may be unsubstituted or mono-, dior trisubstituted by A, OH, OA and/or CN,

A, A' each, independently of one another, denote unbranched, branched or cyclic alkyl having 1-12 C atoms, in which, in addition, 1-7 H atoms may be replaced by F and/or chlorine,

Ar denotes naphthyl, biphenyl, or phenyl which is unsubstituted or mono-, di- or trisubstituted by Hal, A, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³CON(R³)₂, NR³SO₂A, COR³, SO₂N(R³)₂, S(O)_nA, -[C(R³)₂]_n-COOR³ or -O-[C(R³)₂]_p-COOR³,

Hal denotes F, Cl, Br or I,

n denotes 0, 1, 2 or 3, and

p denotes 1, 2, 3, 4 or 5,

including a stereoisomer thereof,

and <u>or a</u> pharmaceutically <u>acceptable salt thereof usable derivatives</u>, <u>solvates</u>, <u>salts and stereoisomers thereof</u>, <u>including mixtures thereof in all ratios</u>.

(Currently Amended) <u>A compound</u> Compounds according to Claim 1 in which

R denotes Hal or -C≡C-H, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

(Currently Amended) <u>A compound</u> Compounds according to Claim 1 in which

R¹ denotes H, =O, Hal, A, OH or OA, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

4. (Currently Amended) <u>A compound</u> Compounds according to claim 1 in which

R¹ denotes OH, and pharmaceutically usable derivatives, solvates, salts and stereoisomers

thereof, including mixtures thereof in all ratios.

- 5. (Cancelled)
- 6. (Currently Amended) <u>A compound</u> Compounds according to claim 1 in which
 - R² denotes H or Hal, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 7. (Withdrawn and Currently Amended) A compound Compounds according to claim 1 in which

 R³ denotes H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 8. (Withdrawn and Currently Amended)

 A compound Compounds

 according to claim 1

 in which
 - Het denotes a monocyclic saturated, unsaturated or aromatic heterocycle having 1 to 2 N and/or O atoms, which may be unsubstituted or mono, di- or trisubstituted by A, OH and/or OA, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- (Withdrawn and Currently Amended) <u>A compound Compounds</u>
 according to claim 1
 in which
 - Het denotes furyl, thienyl, pyrrolyl, imidazolyl, pyridyl, pyrimidinyl, pyrazolyl, thiazolyl, indolyl, pyrrolidinyl, piperidinyl, morpholinyl or piperazinyl, each of which is unsubstituted or mono-, di- or trisubstituted by A, OH and/or OA,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

10. (Withdrawn and Currently Amended)

A compound Compounds
according to claim 1
in which

Het¹ denotes an unsubstituted mono- or bicyclic aromatic heterocycle having 1 to 2 N, O and/or S atoms, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

- 11. (Withdrawn and Currently Amended) A compound Compounds
 according to claim 1
 in which
 R⁵ denotes H or A,
 and pharmaceutically usable derivatives, solvates, salts and stereoisomers
 thereof, including mixtures thereof in all ratios.
- 12. (Withdrawn and Currently Amended)

 A compound Compounds

 according to claim 1

 in which
 - Ar denotes naphthyl, or phenyl which is unsubstituted or mono-, di- or trisubstituted by Hal, A, OR³, N(R³)₂, NO₂, CN, COOR³ or CON(R³)₂, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 13. (Withdrawn and Currently Amended)

 according to claim 1

 in which

Ar denotes phenyl, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

14. (Withdrawn and Currently Amended) <u>A compound Compounds</u>

in which

- R denotes Hal or -C≡C-H,
- R¹ denotes OH,

according to claim 1

- X denotes CH or N,
- X' denotes CH,
- Y denotes R⁴ or Hal,
- R² denotes H or Hal,
- R³ denotes alkyl having 1, 2, 3, 4, 5 or 6 C atoms,
- R⁴ denotes OH, OA, A-COO-, NHA, NHAr, NAA', Het, -NH-CHR⁵-COOR³ or -NH-CHR⁵-COOH,
- R⁵ denotes H or A,
- Het denotes a monocyclic saturated, unsaturated or aromatic heterocycle having 1 to 2 N and/or O atoms, which may be unsubstituted or mono-, di- or trisubstituted by A, OH and/or OA,
- A, A' each, independently of one another, denote unbranched, branched or cyclic alkyl having 1-12 C atoms, in which, in addition, 1-7 H atoms may be replaced by F and/or chlorine,
- Hal denotes F, Cl, Br or I,
- n denotes 0, 1, 2 or 3,
- p denotes 1, 2, 3, 4 or 5,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

15. (Withdrawn and Currently Amended) A compound of Compounds of the formula Ia

according to claim 1

in which

R denotes Hal or -C≡C-H,

R¹ denotes OH,

X denotes CH or N,

X' denotes CH,

Y denotes R⁴ or Hal,

R² denotes H or Hal,

R³ denotes alkyl having 1, 2, 3, 4, 5 or 6 C atoms,

R⁴ denotes OH, OA, A-COO-, NHA, NAA', Het,

-NH-CHR5-COOR3 or -NH-CHR5-COOH,

R⁵ denotes H or A,

Het denotes a monocyclic saturated, unsaturated or aromatic

heterocycle having 1 to 2 N and/or O atoms, which may be

unsubstituted or mono-, di- or trisubstituted by A, OH and/or

OA,

A, A' each, independently of one another, denote unbranched,

branched or cyclic alkyl having 1-12 C atoms, in which, in

addition, 1-7 H atoms may be replaced by F and/or chlorine,

Hal denotes F, Cl, Br or I,

n denotes 0, 1, 2 or 3,

p denotes 1, 2, 3, 4 or 5,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

16. (Withdrawn and Currently Amended) <u>A compound Compounds</u>

- according to Claim 1, which is selected from the group consisting of 1-N-(4-chlorophenyl)-2-N-{4-[(2-dimethylaminoethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(*N*-methyl,*N*-butylamino)-ethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(morpholin-4-yl)ethanoyl)methyl-amino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(4-hydroxypiperidin-1-yl)-ethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(2,6-dimethylmorpholin-4-yl)-ethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(3-cyclohexylmethylpiperidin-1-yl)ethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide.
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-diethylaminoethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(N-methyl,N-ethylamino)ethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-(2-methylimidazol-1-yl)ethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- $1-N-(4-ethynylphenyl)-2-N-\{4-[(2-dimethylaminoethanoyl)methyl-amino]phenyl\}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,$
- 1-N-(4-chlorophenyl)-2-N-{2-fluoro-4-[(2-dimethylaminoethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl) 2-N-{5-[(2-dimethylaminoethanoyl)methyl-amino|pyridin-2-yl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-acetoxyethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- $methyl\ (2R,4R)-2-[(\{[4-(\{1-[1-(4-chlorophenylcarbamoyl)-4-hydroxy-pyrrolidin-2-yl]methanoyl\}amino)phenyl]methylcarbamoyl\}methyl)amino]-4-methylpentanoate,$

- 1-N-(4-chlorophenyl)-2-N-{4-[(2-ethylaminoethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-chloroethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-cyclohexylaminoethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-methylaminoethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-isopropylaminoethanoyl)methyl-amino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-*tert*-butylaminoethanoyl)methyl-amino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-cyclopentylaminoethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-cyclopropylmethylaminoethanoyl)-methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-hydroxyethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-methoxyethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-ethoxyethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-propoxyethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-chlorophenyl)-2-N-{4-[(2-butoxyethanoyl)methylamino]phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 1-N-(4-ethynylphenyl)-2-N-{4-[(2-methoxyethanoyl)methylamino]-phenyl}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, <u>and</u>
- $1-N-(4-chlorophenyl)-2-N-\{2-fluoro-4-[(2-methoxyethanoyl)methyl-amino]phenyl\}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,$
- 1-N (4-chlorophenyl) 2-N {5-[(2-methoxyethanoyl)methylamino]-pyridin-2-yl}-(2R,4R) 4-hydroxypyrrolidine 1,2-dicarboxamide,

and pharmaceutically <u>acceptable</u> usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

- 17. (Withdrawn and Currently Amended)

 A process for preparing a

 compound of claim 1, comprising Process for the preparation of compounds of
 the formula I according to claim 1 and pharmaceutically usable derivatives,
 solvates, salts and stereoisomers thereof, characterised in that
 - a) reacting a compound of the formula II

$$\begin{array}{c|c} R \\ \hline \\ N \\ O \\ R^2 \end{array}$$

in which R, R¹, R², X and X' have the meanings indicated for the compound of formula I in Claim-1,

is reacted with a compound of the formula III

in which Y and R³ have the meanings indicated for the compound of formula I in Claim 1,

or

b) reacting a compound of the formula IV

in which R, R^1 , R^2 , R^3 , X and X' have the meanings indicated for the compound of formula I in Claim 1,

is reacted with a compound of the formula V

in which Y has the meaning indicated <u>for the compound of formula I</u>, in Claim ‡ and L denotes Cl, Br, I or a free or reactively functionally modified OH group,

or

c) reacting a compound of the formula VI

in which R and R¹ have the meanings indicated <u>for the compound of formula I</u> in Claim 1, and L denotes Cl, Br, I or a free or reactively functionally modified OH group,

is reacted with a compound of the formula VII

$$H_2N$$
 X
 R^3
 VII

in which R², R³, X, X' and Y have the meanings indicated for the compound of formula I in Claim 1,

and/or

a base or acid of a compound of the formula I is converted into one of its salts.

- 18. (Currently Amended) Compounds of the formula I according to claim 1 as inhibitors of A method for inhibiting coagulation factor Xa, comprising administering to a subject in need thereof an effective amount of a compound of claim 1.
- 19. (Currently Amended) Compounds of the formula I according to claim 1 as inhibitors of A method for inhibiting coagulation factor VIIa, comprising administering to a subject in need thereof an effective amount of a compound of claim 1.
- 20. (Currently Amended) A pharmaceutical composition comprising a compound Medicaments comprising at least one compound of the formula I according to claim 1 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and optionally excipients and/or adjuvants.
- 21. (Currently Amended) Medicaments comprising at least one compound of the formula I according to claim 1 and/or-pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and

at least one further medicament A pharmaceutical composition according to claim 20, further comprising a further pharmaceutically active ingredient.

- 22. (Withdrawn and Currently Amended) Use of compounds according to claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament for the treatment of A method for treating thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, a tumor, a tumor disease tumours, tumour diseases and/or tumor tumour metastases, comprising administering to a subject in need thereof an effective amount of a compound of claim 1.
- 23. (Currently Amended) A set or kit, comprising Set (kit) consisting of separate packs of
 - (a) an effective amount of a compound of the formula I according to claim 1 and/or a pharmaceutically acceptable salt thereof usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios,

and

- (b) an effective amount of a further <u>pharmaceutically</u> medicament active ingredient.
- 24. (Withdrawn and Currently Amended) Use of compounds of the formula

 I according to claim 1 and/or pharmaceutically usable derivatives, solvates and

 stereoisomers thereof, including mixtures thereof in all ratios,

for the preparation of a medicament for the treatment of A method for treating thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, comprising administering to a subject in need thereof an effective amount of a compound of claim 1 tumours, tumour diseases and/or tumour metastases.

in combination with at least one further medicament active ingredient.